

Amendments to the Claims:

1. (Original) A pyrimidinyl compound

4-[[4-amino-5-bromo-6-(4-cyano-2,6-dimethylphenoxy)-2-pyrimidinyl]amino]benzonitrile, a *N*-oxide, an addition salt, a quaternary amine or a stereochemically isomeric form thereof.

2. (Currently Amended) A pyrimidinyl compound ~~according to claim 1~~ wherein the pyrimidinyl compound is 4-[[4-amino-5-bromo-6-(4-cyano-2,6-dimethylphenoxy)-2-pyrimidinyl]amino]benzonitrile.

3. (Currently Amended) A pharmaceutical composition comprising a pharmaceutically acceptable carrier and ~~a therapeutically active~~ an effective amount of a pyrimidinyl compound according to ~~claims 1 or 2~~ any of claims 1 or 2 or any of claims 23 to 33.

4. (Currently Amended) A combination comprising a pyrimidinyl compound according to ~~claims 1 or 2~~ any of claims 1 or 2 or any of claims 23 to 33 and an antiretroviral compound, wherein said antiretroviral compound comprises at least one of a nucleoside reverse transcriptase inhibitor, a non-nucleoside reverse transcriptase inhibitor, a TIBO compound, an α -APA compound, a TAT-inhibitor, a protease inhibitor, an immunomodulating agent, and mixtures thereof.

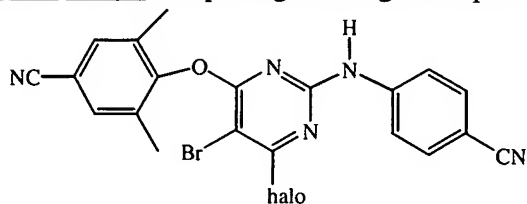
5. (Original) A combination according to claim 4, wherein said nucleoside reverse transcriptase inhibitor comprises at least one of zidovudine (3'-azido-3'-deoxythymidine, AZT), didanosine (dideoxy inosine; ddI), zalcitabine (dideoxycytidine, ddC), lamivudine (3'-thia-2'-3'-dideoxycytidine, 3TC), and mixtures thereof.

6. (Currently Amended) A combination according to claim 4, wherein said non-nucleoside reverse transcriptase inhibitors comprises at least one of suramine, pentamidine, thymopentin, castanospermine, efavirenz, ~~dextran~~ (dextran sulfate), foscarnet-sodium (trisodium phosphono formate), nevirapine (11-cyclopropyl-5,11-dihydro-4-methyl-6*H*-dipyrido[3,2-b : 2',3'-e][1,4]diazepin-6-one), tacrine (tetrahydroaminoacridine), and mixtures thereof.

7. (Original) A combination according to claim 4, wherein said TIBO compound comprises (S)-8-chloro-4,5,6,7-tetrahydro-5-methyl-6-(3-methyl-2-butenyl)imidazo-[4,5,1-jk][1,4]benzodiazepine-2(1*H*)-thione.

8. (Original) A combination according to claim 4, wherein said α -APA compound comprises α -[(2-nitro-phenyl)amino]-2,6-dichlorobenzene-acetamide.
9. (Original) A combination according to claim 4, wherein said protease inhibitor comprises at least one of indinavir, ritanovir, saquinovir, ABT-378, and mixtures thereof.
10. (Original) A combination according to claim 4, comprising at least one of RO-5-3335, levamisole, and mixtures thereof.
11. (Original) A combination according to claim 5, further comprising a pharmaceutically acceptable carrier.
12. (Original) A combination according to claim 6, further comprising a pharmaceutically acceptable carrier.
13. (Original) A combination according to claim 7, further comprising a pharmaceutically acceptable carrier.
14. (Original) A combination according to claim 8, further comprising a pharmaceutically acceptable carrier.
15. (Original) A combination according to claim 9, further comprising a pharmaceutically acceptable carrier.
16. (Original) A combination according to claim 10, further comprising a pharmaceutically acceptable carrier.
17. (Original) A combination according to claim 4 wherein said pyrimidinyl compound and said antiretroviral compound are combined in a single preparation.
18. (Original) A combination according to claim 17, further comprising a pharmaceutically acceptable carrier.

19. (Currently Amended) A process for preparing a compound as claimed in ~~claim 2~~, either of claims 2 or 27, comprising reacting a compound of formula



with NH_3 in the presence of a reaction inert solvent.

20. (Original) A process according to claim 19, wherein said reacting is performed in the presence of a base.

21. (Currently Amended) A method of treating subjects suffering from HIV (Human Immunodeficiency Virus) infection comprising administering to the subject ~~a therapeutically~~ an effective amount of a compound according to ~~claims 1 or 2~~ any of claims 1 or 2 or of claims 23 to 33.

22. (Original) A method of treating subjects suffering from HIV (Human Immunodeficiency Virus) infection comprising administering to the subject a therapeutically effective amount of a combination according to claim 4.

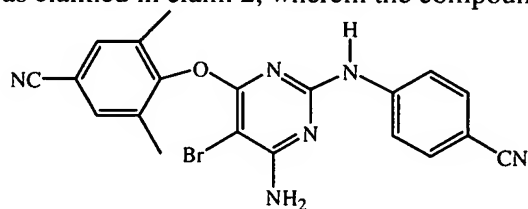
23. (New) A pyrimidinyl compound as claimed in claim 1, wherein the compound is an *N*-oxide of 4-[[4-amino-5-bromo-6-(4-cyano-2,6-dimethylphenyloxy)-2-pyrimidinyl]amino]benzonitrile.

24. (New) A pyrimidinyl compound as claimed in claim 1, wherein the compound is an addition salt of 4-[[4-amino-5-bromo-6-(4-cyano-2,6-dimethylphenyloxy)-2-pyrimidinyl]amino]benzonitrile.

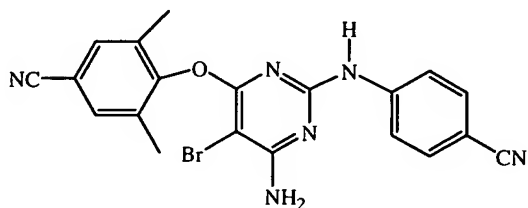
25. (New) A pyrimidinyl compound as claimed in claim 24, wherein the compound is the hydrochloride salt of 4-[[4-amino-5-bromo-6-(4-cyano-2,6-dimethylphenyloxy)-2-pyrimidinyl]amino]benzonitrile.

26. (New) A pyrimidinyl compound as claimed in claim 1, wherein the compound is a quaternary amine of 4-[[4-amino-5-bromo-6-(4-cyano-2,6-dimethylphenyloxy)-2-pyrimidinyl]amino]benzonitrile.

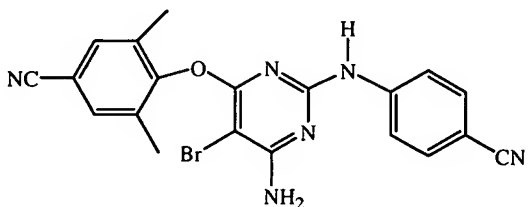
27. (New) A compound as claimed in claim 2, wherein the compound is



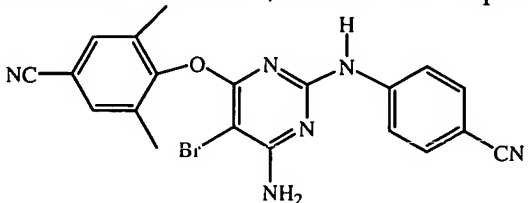
28. (New) A compound as claimed in claim 23, wherein the compound is a N-oxide of



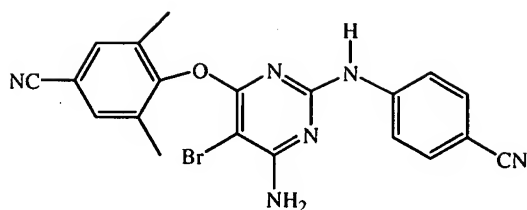
29. (New) A compound as claimed in claim 26, wherein the compound is a quaternary amine of



30. (New) A compound as claimed in claim 24, wherein the compound is an addition salt of



31. (New) A compound as claimed in claim 30, wherein the compound is the hydrochloride salt of



32. (New) An isolated compound as claimed in any of claims 1 or 2 or any of claims 23 to 31.

33. (New) A substantially pure compound as claimed in any of claims 1 or 2 or any of claims 23 to 31.

34. (New) A pharmaceutical composition as claimed in claim 3, wherein the pharmaceutical composition is a tablet.

35. (New) A pharmaceutical composition as claimed in claim 3, wherein the effective amount is between 1 to 1000 mg of active ingredient per unit dosage form.

36. (New) A pharmaceutical composition as claimed in claim 35, wherein the effective amount is between 5 and 200 mg of active ingredient per unit dosage form.

37. (New) A tablet as claimed in claim 34, wherein the effective amount is between 1 to 1000 mg of active ingredient.

38. (New) A tablet as claimed in claim 37, wherein the effective amount is between 5 to 200 mg of active ingredient.

39. (New) A method of treating a subject suffering from HIV-1 (Human Immunodeficiency Virus) infection comprising administering to the subject an effective amount of a compound according to any of claims 1 or 2 or any of claims 23 to 33.

40 (New) A method of treating subjects suffering from HIV-1 (Human Immunodeficiency Virus) that have acquired resistance to art-known non-nucleoside reverse transcriptase inhibitors infection comprising administering to the subject an effective amount of a compound according to any of claims 1 or 2 or any of claims 23 to 33.

41. (New) A combination comprising a pyrimidinyl compound according to any of claims 1 or 2 or any of claims 23 to 33, and an antiretroviral compound, wherein said antiretroviral compound comprises at least one of a nucleoside reverse transcriptase inhibitor, a non-nucleoside reverse transcriptase inhibitor, a TIBO compound, an α -APA compound, a TAT-inhibitor, a protease inhibitor, an immunomodulating agent, and mixtures thereof as a combined preparation for simultaneous, separate or sequential use.

42. (New) A method of treating subjects suffering from HIV (Human Immunodeficiency Virus) infection comprising administering to the subject a compound as claimed in any of claims 1 or 2 or any of claims 23 to 33 and another antiretroviral compound simultaneously, separately or sequentially.

43. (New) A method for inhibiting reverse transcriptase, comprising administering a compound as claimed in any of claims 1 or 2 or any of claims 23 to 33.

44. (New) The method of claim 43, carried out on mammalian cells.

45. (New) The method of claim 43, carried out on human cells.

46. (New) The method of claim 43, carried out on immune cells.

47. (New) The method of claim 43, carried out on human T-4 cells.

48. (New) A complex comprising reverse transcriptase and the compound as claimed in any of claims 1 or 2 or any of claims 23 to 33.